

LOW GAUGE STRETCH WRAP FILM

**[0001]** This utility patent application claims priority to U.S. Provisional Patent Application Serial No. 60/326,834 filed October 3, 2001, entitled Low Gauge Stretch Wrap Film. Serial No. 60/326,834 is hereby incorporated by reference.

Technical Field

**[0002]** The present invention is directed to a multilayer stretch wrap film. More particularly the stretch wrap film of the present invention has a core, a cling layer and a non cling layer.

Background of the Invention

**[0003]** Stretch wrap film is commonly used to wrap a load on a pallet. The film holds goods on a pallet so that the pallet can be stored, shipped and maneuvered. Several physical properties are required for good stretch wrap film. The film should have good stretch, puncture resistance and load containment. The film should also resist tear propagation. The

film must have clarity or good optics so that the contents of the pallet can be viewed through the film.

**[0004]** Stretch wrap film may have a non-cling surface and cling surface opposite the non-cling surface. The non-cling surface generally does not cling to itself and prevents adjoining pallets from sticking to one another. The cling surface enables the film to stick to itself. The cling surface should have sufficient cling so that the film sticks to itself when wrapped around a pallet and does not unravel. However, as cling of the film increases, so does the noise the film creates when it is unwrapped. The noise levels created by unwrapping film with high cling are too high to be tolerated by individuals working around the pallets without ear protection. Therefore, a film is needed with sufficient cling that also yields a tolerable noise level.

**[0005]** There is a constant need to decrease the cost of film for wrapping and shipping pallets. Accordingly, what is needed is a film that has comparable or superior load containment, stretch, puncture resistance and tear resistance to other stretch wrap films but may be used at a lower gauge than commonly used stretch wrap films. The costs of shipping a pallet and the cost of the film to wrap the pallet is also thereby decreased.

#### Summary of the Invention

**[0006]** It is an object of the present invention to provide a stretch wrap film with a balanced set of beneficial properties including high puncture resistance, load containment, stretch, tear resistance and clarity.

**[0007]** Another object of the present invention is to provide a stretch wrap film with sufficient cling that also creates a tolerable noise upon unwrapping.

**[0008]** The films of the present invention have at least three layers, a core layer, a cling layer, and a non-cling layer. Each core layer preferably contains a small percentage of low

density polyethylene combined with linear low density polyethylene, a blend that optimizes several properties, including ultimate stretch, on-pallet stretch, load containment, stretch, puncture resistance and tear resistance of the film. The core layer makes up about 40% to about 96% of the thickness of the film. The film of the present invention may have three or more layers. In a three layer embodiment, the film has a single core layer, a cling layer, and a non cling layer. In a five layer embodiment, the film has two additional core layers. In a seven layer embodiment, the film has five core layers, a cling layer and a non-cling layer.

[0009] The cling layer may contain a balance of plastomer and ultra-low density polyethlyene that optimizes the cling to noise characteristic of the film. The cling layer makes up from about 2% to about 30% of the thickness of the film. In a preferred embodiment the cling layer is about 10% of the overall thickness of the film.

[0010] Additional aspects and advantages of this invention will be apparent from the following detailed description of preferred embodiments thereof, which proceeds with reference to the accompanying drawings.

#### Brief Description of the Drawings

- [0011] FIG. 1 is a depiction of the film of the present invention having five layers.
- [0012] FIG. 2 is a depiction of the film of the present invention having three layers.
- [0013] FIG. 3 is a depiction of the film of the present invention having seven layers.
- [0014] FIG. 4 is a graph illustrating ultimate stretch and on-pallet stretch for Examples 3-9.
- [0015] FIG. 5 is a graph illustrating stretch force and containment force for Examples 3-9.
- [0016] FIG. 6 is a graph illustrating necked-in width, on-pallet puncture energy, and haze at 200% stretch for Examples 3-9.

### Detailed Description of Preferred Embodiments

**[0017]** The present invention relates to stretch films with an optimized balance of beneficial properties. The stretch films of the present invention may include three or more layers, including a core layer, a cling layer and a non-cling layer. Embodiments of the inventive films with five or fewer layers are represented by the construction A/B/C/D/E as shown in FIG. 1. In this construction, A is a cling layer, E is a non-cling layer and B, C, and D may be the same or different polymer resins, selected to optimize the features discussed above in connection with the core layer, namely load containment, stretch, and puncture resistance while maintaining adequate tear propagation.

**[0018]** FIG. 2 depicts an embodiment in which the film has three layers. FIG. 3 depicts an with seven layers, represented by the construction A/B/C<sub>1</sub>/C<sub>2</sub>/C<sub>3</sub>/D/E. The thicknesses shown in these figures are exemplary. For example, the film shown in FIG. 1, has the construction A/B/C/D/E with corresponding percentage thicknesses of 10/15/45/10/20 of the total thickness of the film. The construction A/C/E shown in FIG. 2 is depicted with corresponding percentage thicknesses of 10/70/20. The construction A/ B/C<sub>1</sub>/C<sub>2</sub>/C<sub>3</sub>/D/E, as shown in FIG. 3, is shown with corresponding thicknesses of 10/15/15/15/15/10/20.

**[0019]** Note that various combinations are possible. For example, the film depicted in FIG. 3 may have five identical core layers, B/C<sub>1</sub>/C<sub>2</sub>/C<sub>3</sub>/D. Alternatively, the seven layer film may have four core layers, B/C<sub>1</sub>/C<sub>2</sub>/C<sub>3</sub>, and two adjoining non-cling layers, D/E, and a cling layer A.

**[0020]** The multilayer films of the present invention are typically manufactured by feed block extrusion. Alternatively, the stretch films of the present invention may be made by blown film (tubular) extrusion. Methods for feed block and blown film extrusion are disclosed in The Wiley Encyclopedia of Packaging Technology, pp. 233-38 (Aaron L. Brody

et al. eds., 2<sup>nd</sup> Ed. 1997), which is incorporated herein by reference. Methods for film extrusion are also disclosed in U.S. Pat. No. 6,265,055, which is also incorporated herein by reference.

**[0021]** The cling layer of the film of the present invention enables the film to cling to itself when the film is wrapped on a load. The cling layer is from about 2% to about 30% of the thickness of the film, more preferably about 10%. The cling layer has a balance of ultra low density polyethylene and plastomer enabling the film to have sufficient cling without creating an intolerable noise level upon unwrapping than film.

**[0022]** The cling layer may contain from about 60% to 100% by weight of an ultra low density polyethylene. Ultra low density polyethylene (hereinafter “ULDPE”) is sometimes also known as very low density polyethylene. ULDPE is an ethylene copolymerized with at least one C<sub>3</sub>-C<sub>10</sub> α-olefin, more preferably a C<sub>8</sub> α-olefin. The resulting ULDPE preferably has a density ranging from 0.888 g/cm<sup>3</sup> to 0.920 g/cm<sup>3</sup>, more preferably from about 0.885 g/cm<sup>3</sup> to about 0.920 g/cm<sup>3</sup>, and a melt index ranging from 1.0 g/10 min to 20 g/10 min, more preferably ranging from about 2 g/10 min to about 10 g/10 min. In a preferred embodiment, the cling layer contains about 85% of a ULDPE that is a polyethylene copolymerized with a C<sub>8</sub> α-olefin with a density of about 0.900g/cm<sup>3</sup>, and a melt index of about 5.

**[0023]** ULDPE can be produced by a variety of processes, including gas phase, solution and slurry polymerization as disclosed in The Wiley Encyclopedia of Packaging Technology pp. 748-50 (Aaron L. Brody et al. eds., 2<sup>nd</sup> Ed. 1997), referenced above. ULDPE is typically manufactured using a Ziegler-Natta catalyst, although a number of other catalysts may be used. For example, ULDPE may be manufactured with a metallocene catalyst. Alternatively, ULDPE may be manufactured with a catalyst that is a hybrid of a metallocene catalyst and a Ziegler-Natta catalyst. Methods for manufacturing ULDPE are also disclosed in U.S. Pat. No. 5,399,426, U.S. Pat. No. 4,668,752, U.S. Pat. No. 3,058,963, U.S. Pat. No.

2,905,645, U.S. Pat. No. 2,862,917, U.S. Pat. No. 2,699,457, which are incorporated by reference herein. The density of ULDPE is achieved by copolymerizing ethylene with a sufficient amount of one or more monomers. The monomers are preferably selected from butene, hexene, 4-methyl-1-pentene and octene.

**[0024]** The cling layer may also include a plastomer in a range from 0% to about 40% by weight of the cling layer, more preferably about 15%. The plastomer is preferably a polyethylene copolymerized with a C<sub>3</sub>-C<sub>10</sub> α-olefin, having a density in the range from about 0.850 g/cm<sup>3</sup> to about 0.900 g/cm<sup>3</sup>, more preferably of about 0.860 g/cm<sup>3</sup> to about 0.880 g/cm<sup>3</sup>, and a melt index of about 1.0 g/10 min to about 20.0 g/10 min, more preferably from about 2.0 to about 10.0 g/10 min. The cling layer preferably contains 15% plastomer with a density of about 0.875g/cm<sup>3</sup> and a melt index of about 3.0g/10 min. The plastomer is manufactured by the same process described above with reference to ULDPE, except that a higher amount by weight of one or more monomer is copolymerized with polyethylene. In a preferred embodiment, the plastomer is a polyethylene copolymerized with a C<sub>8</sub> α-olefin, which has an unstretched cling of 250 g and a 200% stretched cling of 66 g as measured by ASTM D 4649. Cling is the strength required, in grams, to pull apart overlapping film along a test section.

**[0025]** While ULDPE provides the film with limited cling, the plastomer provides greater cling in the resulting film. Films of the present invention containing percentage weight of ULDPE in the higher ranges specified herein for ULDPE create less noise during unwrapping and have less cling. In contrast, films with a higher percentage weight of plastomer provide greater cling but more noise. The ranges provided for ULDPE and plastomer provide a balanced film, with a sufficient amount of cling and an intermediate noise level.

**[0026]** The stretch film of the present invention has one or more core layers. The core layers or layer make up about 40% to about 96% of the thickness of the film, more preferably

about 70% of the thickness of the film. Each core layer includes a low density polyethylene in an amount ranging from about 0.01% to about 3% by weight of the core layer and a linear low density polyethylene (hereinafter "LLDPE") in an amount ranging from about 97 to about 99.9% by weight of the core layer. More particularly, the core layer includes a low density polyethylene in an amount less than 3%.

**[0027]** The LDPE of the core layer may have a density ranging from about 0.900 g/cm<sup>3</sup> to about 0.940 g/cm<sup>3</sup>, more preferably from about 0.920 g/cm<sup>3</sup> to about 0.930 g/cm<sup>3</sup> and a melt index ranging from about 0.10 g/10 min to about 10.0 g/10 min, more preferably from about 0.2 g/10 min to about 0.4 g/10 min. LDPE is generally used for heavy bags, such as ice bags or shipping sacks, which normally have a thickness from 1.5 mm to 4.0 mm. In contrast, the film of the present invention preferably has a thickness of 0.5 to 1.5 mm.

**[0028]** The LDPE may be an ethylene homopolymer or an ethylene copolymerized with one or monomer, such as vinyl acetate or a C<sub>3</sub>-C<sub>10</sub> α-olefin. The LDPE is preferably ethylene homopolymer with a density of about 0.921 g/cm<sup>3</sup> and a melt index of about 0.2 g/10 min. An alternative embodiment of the stretch film contains an LDPE that is a copolymer. Methods for manufacturing LDPE are disclosed in The Wiley Encyclopedia of Packaging Technology (Aaron L. Brody et al. eds., 2<sup>nd</sup> Ed. 1997) pp. 753-754 and in U.S. Pat. No. 5,399,426, both of which are referenced above.

**[0029]** The LLDPE is a polyethylene copolymerized with one or more C<sub>3</sub>-C<sub>10</sub> α-olefins, with a density ranging from about 0.900 to about 0.940 g/cm<sup>3</sup>, more preferably from about 0.910 to about 0.930 g/cm<sup>3</sup>, and a melt index ranging from about 1.0 to about 20.0 g/10 min., more preferably from about 1.5 to about 6.0 g/10 min. Similar to the ULDPE discussed above, LLDPE used in the films of the present invention is preferably an ethylene copolymerized with one of butene, hexene, 4-methyl-1-pentene, or octene. LLDPE may also be polymerized with various copolymers of butene, hexene, 4-methyl-1-pentene, and/or

octene. The core layer preferably contains about 97% to about 99.9% of a LLDPE that is an ethylene copolymerized with a C<sub>8</sub> α-olefin, with a density of 0.917 g/cm<sup>3</sup> and a melt index of 4.0 g/10 min.

**[0030]** LLDPE is manufactured by the same processes discussed above in connection with ULDPE. For example, ethylene may be copolymerized using various catalysts such as a Ziegler-Natta catalyst or a metallocene catalyst. LLDPE has a higher density than ULDPE and plastomer because it is copolymerized with a lower concentration of comonomer than ULDPE or plastomer.

**[0031]** The density ranges and melt indexes as disclosed herein for ULDPE and LLDPE and for plastomer and ULDPE partially overlap. However, these polymers may be further distinguished by crystallinity, measured as melt point. A ULDPE with the same melt index and density of LLDPE will have a lower melt point than LLDPE as disclosed herein. Similarly, a plastomer with the same density and melt index of a ULDPE, will have a lower melt point than ULDPE.

**[0032]** As mentioned above, the film of the present invention further includes a non-cling layer of polypropylene. The polypropylene non-cling layer may have a density ranging from about 0.890 g/cm<sup>3</sup> to about 0.910 g/cm<sup>3</sup>, more preferably from about 0.895 g/cm<sup>3</sup> to about 0.905 g/cm<sup>3</sup> and a melt flow rate from about 2.0 g/10 min to about 40.0 g/10 min. The polypropylene may be a homopolymer or may be copolymerized with a comonomer, preferably ethylene. Alternatively, polypropylene may be copolymerized with another comonomer, such as a C<sub>3</sub>-C<sub>10</sub> α-olefin. The polypropylene of the present invention is preferably a polypropylene copolymer and ethylene in an amount ranging from 0 to about 10% by weight of the polypropylene, more preferably in an amount ranging from about 1 to about 5% by weight of the polypropylene. A preferred embodiment has a polypropylene copolymerized with ethylene that is included in an amount ranging from about 2% to about

4% by weight of the polypropylene, known as random copolymer polypropylene. The random copolymer of polypropylene may have a density of about 0.905 g/cm<sup>3</sup> and a melt flow rate of about 10.0 g/10 min. The method for measuring polypropylene melt flow rate is disclosed in The Wiley Encyclopedia of Packaging Technology (Aaron L. Brody et al. eds., 2<sup>nd</sup> Ed. 1997) p. 677 and methods for manufacturing polypropylene are disclosed in Kirk-Othmer Concise Encyclopedia of Chemical Technology pp. 1420-21 (Jacqueline I. Kroschwitz et al. eds., 4<sup>th</sup> Ed. 1999), which is incorporated herein by reference.

**[0033]** The amount of each polymer included in the inventive films is selected to create an optimal balance of several physical properties. The stretch films of the present invention balance increased load and holding containment and stretch with clarity, puncture resistance and resistance to tear propagation. The inventive stretch films have a core layer that includes linear low density polyethylene with a small percentage of low density polyethylene. The low density polyethylene provides good load containment, stretch and necked-in width. As will be discussed in more detail below in conjunction with examples, the amount of low density polyethylene is a low percentage, in the range of 0.01% to about 3%, because at higher percentages, low density polyethylene reduces the film's puncture resistance, resistance to tear propagation and clarity. Therefore, the stretch film of the present invention optimizes load containment, stretch, puncture resistance and clarity by incorporating a small percentage of low density polyethylene. that is no greater than about 90 gauge

**[0034]** It has been discovered that the greatest benefits from the balance of properties of the present invention are found in gauges ranging from about 45 gauge to about 90 gauge. Even more beneficial are the properties of the film of the present invention at gauges ranging from about 50 gauge to about 80 gauge. At these gauge ranges, it has been discovered that the film performs better than a higher gauge film of the prior art, in terms of load containment, stretch, puncture resistance, clarity and tear propagation resistance. Therefore,

the films of the present invention preferably have a gauge ranging from about 45 gauge to about 90 gauge, more preferably from about 50 gauge to about 80 gauge. At gauges higher than about 90, the films of the present invention do not perform significantly better in terms of load containment, stretch, puncture resistance, clarity and tear propagation resistance when compared with prior art films having the same gauge. Note, however, that films having gauges higher than about 90 gauge still perform satisfactorily in terms of of load containment, stretch, puncture resistance, clarity and tear propagation resistance. Note also that these films of the present invention are designed so that they have sufficient cling while also creating tolerable noise upon unwrapping.

**[0035]** It is believed that the enhanced performance of the inventive films is optimized at these gauges because of the manner in which the films are used. Generally, stretch wrap machines run film at a pre-stretch of 200% to 300%. At this level of pre-stretch, the inventive films below a gauge of about 90 will provide higher resistance to stretch and become stiffer than films of gauges above about 90. The increased level of stiffness at a pre-stretch of 200% to 300% provides enhanced load containment and puncture resistance for inventive films having gauges no greater than about 90 gauge or 80 gauge.

**[0036]** Because of the beneficial properties of the inventive films at low gauges, the optimized balance of stretch, load containment, puncture resistance and clarity, the films may be used at a decreased gauge or thickness over stretch films of the prior art. Conventionally, stretch films range from about 0.4 mm to about 3 mm. Because of the balanced properties of the film of this invention, a lower gauge film can be used while achieving superior performance as compared to prior art films of higher gauges. As a result, the end user can use less weight of the inventive film to wrap a pallet than would be required to wrap the same load with a film of the prior art. Since less film is used, it provides a cost advantage.

**[0037]** One skilled in the art will recognize that the overall thickness of the stretch film can be varied depending on the particular end use for which the film is manufactured. The films of the present invention have a thickness that is generally in the range of typical thickness for stretch wrap films, but may be easily adjusted by one skilled in the art to fit the desired end use. However, as discussed above, the beneficial properties of the films of the present invention, such as increased stretch puncture resistance and load containment, may allow the films to be used at a decreased gauge or thickness over stretch films of the prior art.

**[0038]** In addition to the properties discussed above, the stretch films of the present invention optimize the amount of cling while minimizing the noise created upon unwrapping film. The noise of the film unwrapping increases as the cling of the film increases. In the cling layer, the amount of ultra low density polyethylene, a polymer that provides intermediate cling, is balanced against the amount of plastomer, a lower density polyethylene with high cling. As a result, the film has a cling layer that provides sufficient cling while creating tolerable noise upon wrapping or unwrapping.

### **EXAMPLES OF THE PREFERRED EMBODIMENTS**

**[0039]** Examples are provided of the present invention in order to compare the properties of the composition by varying the content of the layers in the films. Example 1 describes a stretch film composition prepared in accordance with the present invention. Example 2 is a hypothetical example of the inventive film. Examples 3-9 describe film compositions similar to Example 1, except that the level of LDPE in the core layer is varied from 0% to 6% incrementally by one percentage. Tables 3-9 describe the physical properties of the Examples 3-9 which are graphically presented in FIGS. 3-6. Comparative Examples 1-28 compare a film as described in Example 1 with a prior art film.

## Example 1

[0040] Example 1 is a film containing five layers, as represented by the construction A/B/C/D/E with corresponding percentage thicknesses of 10/15/45/10/20 of the total thickness of the film. The polymers in the film as described in the table below are available from the Dow Chemical Company (hereinafter "Dow") in Houston, Texas. The Dow trade name of each polymer is listed in parenthesis next to its description. The film was formed from the following components:

**Table A**

<b>Layer</b>	<b>Composition</b>	<b>% of Layer by Wt.</b>
A	Polyethylene copolymerized with a C <sub>8</sub> olefin, with a density of 0.900 g/cm <sup>3</sup> and a melt index of 5.0g/10 min. (ULDPE) (61520.09L) Polyethylene copolymerized with a C <sub>8</sub> olefin, with a density of 0.875 g/cm <sup>3</sup> and a melt index of 3.0 g/10 min (Plastomer) (KC8852)	85% 15%
B, C, D	Polyethylene copolymerized with a C <sub>8</sub> olefin, with a density of 0.917 g/cm <sup>3</sup> and a melt index of 4.0 g/10 min. (LLDPE) (Elite 5230) Polyethylene homopolymer with a density of g/cm <sup>3</sup> and a melt index of 0.2 g/10 min. (LDPE) (LDPE 132i)	97% 3%
E	Random copolymer polypropylene with a density of 0.905 g/cm <sup>3</sup> and a melt flow rate of 10.0 g/10 min. (13510A)	100%

[0041] The film, with a 29.2 micron thickness (1.15 gauge), had the following properties:

- (a) high puncture resistance, about 180 g/mil, by dart drop impact F-50 (ASTM D1709);
- (b) good optical quality, less than 3% haze (ASTM D1003);
- (c) high load containment and strength, 5.9 KPSI in the machine direction and 4.4 KPSI in the cross direction (ASTM D882);
- (d) good stretch, 540% in the machine direction and 640% in the cross direction (ASTM D882);
- (e) good tear resistance, 390 g/mil MD, and 690 g/mil CD (ASTM D1922);

and (f) sufficient non-cling on the non-cling side of the film, coefficient of friction of 1.8 (ASTM D1894). The film had a cling of about 190 grams unstretched and a cling of about 150 grams at 200% stretch according to ASTM test method D4649. As discussed above, the film of the present invention also has a tolerable noise level while still providing sufficient cling.

### Example 2

[0042] In another embodiment, the film has three layers, represented by the construction A/C/E, wherein the layers have a corresponding percentages of thickness of 10/70/20. Such a film is illustrated in FIG. 2. The polymers in the film may be obtained, as specified below, from Dow in Houston, Texas, ExxonMobil Chemical (“ExxonMobil”) in Houston, Texas and Equistar Chemical Company (“Equistar”) in Houston, Texas. The film has the following composition:

TABLE B

Layer	Composition	% of Layer by Wt.
A	Polyethylene copolymerized with a C <sub>6</sub> α-olefin, with a density of 0.900 g/cm <sup>3</sup> and a melt index of 5.0g/10 min. (ULDPE) (Dow, formerly Union Carbide, Flexomer 9042)	85%
	Polyethylene copolymerized with a C <sub>4</sub> α-olefin with a density of 0.873 g/cm <sup>3</sup> and a melt index of 4.5 g/10 min. (plastomer) (ExxonMobil Exact 4049)	15%
C	Polyethylene copolymerized with a C <sub>6</sub> α-olefin with a density of 0.917 g/cm <sup>3</sup> and a melt index of 3.5 g/10 min. (LLDPE) (ExxonMobil Exceed 357C32)	97%
	Polyethylene homopolymer with a density of 0.918 g/cm <sup>3</sup> and a melt index of 0.25 g/10 min. (LDPE) (Equistar Petrothene NA 940)	3%

E Random copolymer polypropylene with a density 100% of 0.900 g/cm<sup>3</sup> and a melt flow rate of 10.0 g/10 min. (Equistar Petrothene PP8310-GO)

[0043] This film would be expected to have optimized beneficial properties, including good cling but tolerable noise, good tear resistance, high puncture resistance, high load containment and high stretch capabilities. As will be recognized by one skilled in the art, although Example 1 identifies a five layer film and Example 2 identifies a three layer film, the same compositions identified can be used to make a stretch wrap film which includes one or more core layers, at least one cling layer and at least one non-cling layer. For example, the same compositions may be used to make a seven layer film with five core layers, one cling layer and one non-cling layer.

### Examples 3-9

[0044] Seven examples, Examples 3-9, have been tested with varying levels of LDPE. Examples 3-9 contain the same composition as described above in Example 1 with two variations. In Example 3, the core layer contains no LDPE. In each remaining comparative example, the core layer contains a small percentage of LDPE. Example 4 contains a core layer with 1% LDPE; Example 5 contains a core layer with 2% LDPE; Example 6 contains a core layer with 3% LDPE; Example 7 contains a core layer with 4% LDPE; Example 8 contains a core layer with 5% LDPE; and Example 9 contains a core layer with 6% LDPE. In addition, each of the six examples contains a polypropylene resin that is designated Dow DS6D82 instead of the Dow 13510A that is identified in Example 1. Dow DS6D82 is a random copolymer polypropylene resin with a density of 0.9 g/cm<sup>3</sup> and a melt flow rate of 7 g/10 min.

**[0045]** These examples were tested for the following seven properties: ultimate stretch, on-pallet stretch, stretch force at 100%, containment force, necked-in width, on-pallet puncture energy (puncture resistance) and haze at 200% stretch. Ultimate stretch is tested by incrementally stretching the film from 0% until the film breaks and measuring the stretch percentage at which the film breaks. Stretch force is tested by meausing the amount of ofrce necessary to stretch the film to a specifiedid percentage, 100% in this case. Ultimate stretch and stretch force represent measurements of the film's elongational properties. Haze was measured by ASTM D1003, a method that includes stretching the film to 200% and testing the haze with a hazemeter.

**[0046]** On-pallet stretch, necked-in width, containment force and puncture energy are all conducted with a Lantech Q300, a turntable stretch wrapper. The results for these comparative examples were obtained by running the Lantech Q300 at 200% prestretch with 16 lbs of secondary force and 9 wraps around a testing cage. On-pallet stretch is measured by marking the roll as it unwinds and then measuring the distance between the marks on the cage. Necked-in width is the measurement of the width of the film on the cage as compared to the original width, which is 20 inches. Containment force is a measurement of the force that 9 wraps of film exert on a load cell bar that is attached to the cage. Puncture energy is measured with a hand force gauge that is punched through the film. The hand force gauge measure the amount of force required to punch through the film.

**[0047]** Table C shows the ultimate stretch for each of the six comparative examples. Table D shows the on-pallet stretch for each example. The data contained in these tables is illustrated in FIG. 4. Table C shows the stretch force at 100% for each example and Table E shows the containment force for each examples. The data of Tables E and F is illustrated in FIG. 5. Table G shows necked in width, Table H shows on-pallet puncture energy and Table I shows haze at 200% stretch. FIG. 6 illustrates these three properties in graph form.

**TABLE C****Ultimate Stretch (%)**

<b>Example</b>	<b>Test 1</b>	<b>Test 2</b>	<b>Test 3</b>	<b>Ave</b>	<b>Std. Deviation</b>
3	305	308	308	307.00	1.73
4	336	325	339	333.33	7.37
5	360	339	341	346.67	11.59
6	349	336	335	340.00	7.81
7	370	364	363	365.67	3.79
8	353	360	361	358.00	4.36
9	351	314	359	355.00	5.66

**TABLE D****On Pallet Stretch**

<b>Example</b>	<b>Stretch (%)</b>
3	222.5
4	217.5
5	222.5
6	212.5
7	207.5
8	212.5
9	202.5

**TABLE E****Stretch Force at 100% Stretch (lbs.)**

<b>Example</b>	<b>Test 1</b>	<b>Test 2</b>	<b>Test 3</b>	<b>Average</b>	<b>Std. Deviation</b>
3	38	38	40	38.67	1.15
4	38	38	37	37.67	0.58
5	41	40	40	40.33	0.58
6	42	42	42	42.00	0
7	44	45	45	4467	0.58
8	45	45	48	46.00	1.73
9	46	45	46	45.67	0.58

**TABLE F**

**Containment Force**

<b>Example</b>	<b>Force (lbs.)</b>
3	31.5
4	31.5
5	30.5
6	31.5
7	33.0
8	33.0
9	34

**TABLE G**

**Stretch Force at 200% Haze (%)**

<b>Example</b>	<b>Test 1</b>	<b>Test 2</b>	<b>Test 3</b>	<b>Test 4</b>	<b>Test 5</b>	<b>Average</b>	<b>Std. Deviation</b>
3	0.75	0.79	0.79	0.79	0.84	0.79	0.031937
4	0.75	0.82	1.03	0.82	0.69	0.82	0.128335
5	0.91	1.05	1.06	1.1	0.95	1.01	0.080187
6	1.29	1.22	1.23	1.29	1.26	1.26	0.032711
7	1.21	1.15	1.06	1.16	1.21	1.16	0.0614
8	1.27	1.36	1.67	1.5	1.49	1.46	0.152217
9	1.32	1.62	1.63	1.9	1.59	1.61	0.205597

**TABLE H**

**Necked-In Width Based on 20 Inch Film**

<b>Example</b>	<b>Width (inches)</b>
3	15.00
4	15.25
5	15.25
6	15.75
7	16.00
8	15.75
9	16.25

**TABLE I**  
**On Pallet Puncture Energy**

<b>Example</b>	<b>lbs.</b>
3	9.50
4	9.25
5	9.25
6	9.25
7	8.75
8	8.50
9	9.00

**[0048]** As can be seen by Tables C-F and FIGS. 4-5, ultimate stretch, stretch force and containment force rise as the level of LDPE in the core layer is increased above 3%. Similarly, as can be seen in Table G and FIG. 6, necked-in width increases slightly at the level of LDPE in the core layer reaches 4%. However, as can be seen in Tables H-I and FIG. 6, haze and on-pallet puncture energy decrease as the level of LDPE in the core layer is increase above 3%. At a level of about 3% or less LDPE in the core layer, the film has good stretch, strength and necked-in width while having optimal clarity and puncture resistance. Therefore, levels of LDPE in the core layer at about 3% or below provide an optimal balance of the beneficial properties provided by LDPE.

### **Comparative Examples 1-28**

**[0049]** Testing for the following examples was performed with a variety of different equipment and at a variety of settings. Some of the tests were performed with a Lantech Q300, some with Orion equipment and others with Signode equipment. Each of the examples discussed below compares a film prepared from the composition detailed in Example 1 (referred to hereafter as “Sample Film”) to a film of the prior art. However, since the testing was conducted on different equipment, films prepared from the composition in Example 1 having the same gauge produced different results.

### **Comparative Example 1**

**[0050]** A film was prepared corresponding with the composition detailed in Example 1 for comparison with a film manufactured by AEP under the tradename A12090. The AEP film was a 90 gauge film while the film of Example 1 was a 63 gauge film. It was determined that the AEP film had a weight/load of 9.9 ounces and the Example 1 film had a film weight/load of 4.9 ounces. The AEP film had a containment force of 23 lbs compared with 28 lbs for the film of Example 1.

### **Comparative Example 2**

**[0051]** A Sample Film was prepared for comparison with a film manufactured by AEP under the tradename A1 2090. The AEP film was a 90 gauge film while the Sample Film was a 51 gauge film. It was determined that the AEP film had a film weight/load of 12.9 ounces and the Sample Film had a film weight/load of 5.2 ounces.

### **Comparative Example 3**

**[0052]** A Sample Film was prepared for comparison with a film manufactured by Tyco/Armin. The Tyco/Armin film was a 70 gauge film while the Sample Film was a 51 gauge film. It was determined that the Tyco/Armin film had a film weight/load of 3.1 ounces and the Sample Film had a film weight/load of 1.7 ounces. The Tyco/Armin film had a containment force of 23 lbs compared with 25 lbs for the Sample Film.

### **Comparative Example 4**

**[0053]** Two Sample Films were prepared for comparison with a film manufactured by Linear under the tradename ADU26720. The Linear film was a 105 gauge film while the first Sample Film was a 71 gauge film and the second Sample Film was a 105 gauge. It was determined that the Linear film had a film weight/load of 14.4 ounces. The 71 gauge Sample Film had a film weight/load of 9.8 ounces and the 105 gauge Sample Film had a film

weight/load of 14.4 ounces. The Linear film had a containment force of 51 lbs compared with 46 lbs for the 71 gauge Sample Film and 45 lbs for the 105 gauge Sample Film.

#### **Comparative Example 5**

**[0054]** A Sample Film was prepared for comparison with a film manufactured by AEP. The AEP film was an 80 gauge film while the Sample Film was a 51 gauge film. It was determined that the AEP film had a film weight/load of 5.3 ounces and the Sample Film had a film weight/load of 3.3 ounces. The AEP film had a containment force of 23.5 lbs compared with 27 lbs for the Sample Film.

#### **Comparative Example 6**

**[0055]** A Sample Film was prepared for comparison with a film manufactured by Tyco under the tradename AMP2. The Tyco film was a 70 gauge film while the Sample Film was a 51 gauge film. It was determined that the Tyco film had a film weight/load of 8.3 ounces and the Sample Film had a film weight/load of 4.9 ounces. The Tyco film had a containment force of 26 lbs compared with 26 lbs for the Sample Film.

#### **Comparative Example 7**

**[0056]** A Sample Film was prepared for comparison with a film manufactured by Linear under the tradename ADC 17820V. The Linear film was a 70 gauge film while the Sample Film was a 51 gauge film. It was determined that the Linear film had a film weight/load of 6.4 ounces and the Sample Film had a film weight/load of 4.5 ounces. The Linear film had a containment force of 25 lbs compared with 25 lbs for the Sample Film.

#### **Comparative Example 8**

**[0057]** A Sample Film was prepared for comparison with a film manufactured by Amtopp. The Amtopp film was an 80 gauge film while the Sample Film was a 51 gauge film. It was determined that the Amtopp film had a film weight/load of 5.3 ounces and the

Sample Film had a film weight/load of 3.3 ounces. The Amtopp film had a containment force of 20.5 lbs compared with 27 lbs for the Sample Film.

#### **Comparative Example 9**

**[0058]** A Sample Film was prepared for comparison with a film manufactured by Sigma. The Sigma film was a 100 gauge film while the Sample Film was a 71 gauge film. It was determined that the Sigma film had a film weight/load of 5.1 ounces and the Sample Film had a film weight/load of 3.1 ounces. The Sigma film had a containment force of 48.5 lbs compared with 47.5 lbs for the Sample Film.

#### **Comparative Example 10**

**[0059]** A Sample Film was prepared for comparison with a film manufactured by AEP under the tradename TLC. The AEP film was an 80 gauge film while the Sample Film was a 51 gauge film. It was determined that the AEP film had a film weight/load of 5.5 ounces and the Sample Film had a film weight/load of 2.8 ounces. The AEP film had a containment force of 21.5 lbs compared with 23.0 lbs for the Sample Film.

#### **Comparative Example 11**

**[0060]** A Sample Film was prepared for comparison with a film manufactured by Linear under the tradename ADC8030V. The AEP film was an 80 gauge film while the Sample Film was a 63 gauge film. It was determined that the AEP film had a film weight/load of 28.7 ounces and the Sample Film had a film weight/load of 11.5 ounces. The AEP film had a containment force of 36 lbs compared with 43 lbs for the Sample Film.

#### **Comparative Example 12**

**[0061]** A Sample Film was prepared for comparison with a film manufactured by Paragon. The Paragon film was a 75 gauge film while the Sample Film was a 51 gauge film. It was determined that the Paragon film had a film weight/load of 7.7 ounces and the Sample

Film had a film weight/load of 5.3 ounces. The Paragon film had a containment force of 17 lbs compared with 21 lbs for the Sample Film.

#### **Comparative Example 13**

**[0062]** A Sample Film was prepared for comparison with a film manufactured by AEP under the tradename PP1 8020. The AEP film was an 80 gauge film while the Sample Film was a 51 gauge film. It was determined that the AEP film had a film weight/load of 17.6 ounces and the Sample Film had a film weight/load of 8.3 ounces. The AEP film had a containment force of 19 lbs compared with 22 lbs for the Sample Film.

#### **Comparative Example 14**

**[0063]** A Sample Film was prepared for comparison with a film manufactured by Tyco. The Tyco film was a 70 gauge film while the Sample Film was a 51 gauge film. It was determined that the Tyco film had a film weight/load of 4.0 ounces and the Sample Film had a film weight/load of 2.6 ounces. The Tyco film had a containment force of 26 lbs compared with 28 lbs for the Sample Film.

#### **Comparative Example 15**

**[0064]** A Sample Film was prepared for comparison with a film manufactured by Tyco under the tradename MP2. The Tyco film was a 70 gauge film while the Sample Film was a 51 gauge film. It was determined that the Tyco film had a film weight/load of 7.6 ounces and the Sample Film had a film weight/load of 4.0 ounces. The Tyco film had a containment force of 16 lbs compared with 17 lbs for the Sample Film.

#### **Comparative Example 16**

**[0065]** A Sample Film was prepared for comparison with a film manufactured by Tyco. The Tyco film was an 80 gauge film while the Sample Film was a 51 gauge film. It was determined that the Tyco film had a film weight/load of 4.8 ounces and the Sample Film had

a film weight/load of 2.9 ounces. The Tyco film had a containment force of 37 lbs compared with 34.5 lbs for the Sample Film.

#### **Comparative Example 17**

**[0066]** A Sample Film was prepared for comparison with a film manufactured by AEP under the tradename A1. The AEP film was a 75 gauge film while the Sample Film was a 51 gauge film. It was determined that the AEP film had a film weight/load of 2.4 ounces and the Sample Film had a film weight/load of 1.5 ounces. The AEP film had a containment force of 26 lbs compared with 28 lbs for the Sample Film.

#### **Comparative Example 18**

**[0067]** Two Sample Films were prepared for comparison with a film manufactured by Tyco. The Tyco film was an 80 gauge film while the first Sample Film was a 51 gauge film and the second Sample Film was a 63 gauge film. It was determined that the Tyco film had a film weight/load of 8.5 ounces. The 51 gauge Sample Film had a film weight/load of 4.1 ounces and the 63 gauge Sample Film had a film weight/load of 5.6 ounces.

#### **Comparative Example 19**

**[0068]** A Sample Film was prepared for comparison with a film manufactured by Linear under the tradename ADC-7020. The AEP film was a 70 gauge film while the Sample Film was a 51 gauge film. It was determined that the AEP film had a film weight/load of 5.06 ounces and the Sample Film had a film weight/load of 3.6 ounces. The AEP film had a containment force of 19 lbs compared with 16 lbs for the Sample Film.

#### **Comparative Example 20**

**[0069]** A Sample Film was prepared for comparison with a film manufactured by Linear under the tradename ADC-7020. The Linear film was an 80 gauge film while the Sample Film was a 51 gauge film. It was determined that the Linear film had a film weight/load of

7.9 ounces and the Sample Film had a film weight/load of 4.8 ounces. The Linear film had a containment force of 15 lbs compared with 22 lbs for the Sample Film.

#### **Comparative Example 21**

**[0070]** Two Sample Films were prepared for comparison with a film manufactured by Pactiv under the tradename QM1 2010. The Pactiv film was a 90 gauge film while the first Sample Film was an 80 gauge film and the second Sample Film was a 71 gauge film. It was determined that the Pactiv film had a film weight/load of 7.7 ounces. The 80 gauge Sample Film had a film weight/load of 5.4 ounces and the 71 gauge Sample Film had a film weight/load of 5.0 ounces. The Pactiv film had a containment force of 38 lbs compared with 36 lbs for the 80 gauge Sample Film and 42 lbs for the 71 gauge Sample Film.

#### **Comparative Example 22**

**[0071]** A Sample Film was prepared for comparison with a film manufactured by Sigma under the tradename Select Cast. The Sigma film was an 80 gauge film while the Sample Film was a 63 gauge film. It was determined that the Sigma film had a film weight/load of 4.8 ounces and the Sample Film had a film weight/load of 4.1 ounces. The Sigma film had a containment force of 17 lbs compared with 22 lbs for the Sample Film.

#### **Comparative Example 23**

**[0072]** A Sample Film was prepared for comparison with a film manufactured by Paragon. The Paragon film was a 70 gauge film while the Sample Film was a 51 gauge film. It was determined that the Paragon film had a film weight/load of 6.2 ounces and the Sample Film had a film weight/load of 4.9 ounces. The Paragon film had a containment force of 33 lbs compared with 35 lbs for the Sample Film.

#### **Comparative Example 24**

**[0073]** A Sample Film was prepared for comparison with a film manufactured by Pactiv under the tradename MP2. The Pactiv film was a 100 gauge film while the Sample Film was

a 71 gauge film. It was determined that the Pactiv film had a film weight/load of 11.6 ounces and the Sample Film had a film weight/load of 8.1 ounces.

#### **Comparative Example 25**

[0074] A Sample Film was prepared for comparison with a film manufactured by AEP under the tradename TLC. The Pactiv film was a 70 gauge film while the Sample Film was a 51 gauge film. It was determined that the Pactiv film had a film weight/load of 8.7 ounces and the Sample Film had a film weight/load of 5.5 ounces. The Pactiv film had a containment force of 25 lbs compared with 35 lbs for the Sample Film.

#### **Comparative Example 26**

[0075] Two Sample Films were prepared for comparison with a film manufactured by Paragon under the tradename G2.07520N.50. The Paragon film was a 75 gauge film while the first Sample Film was a 63 gauge film and the second Sample Film was a 51 gauge film. It was determined that the Paragon film had a film weight/load of 17.5 ounces. The 63 gauge Sample Film had a film weight/load of 14.5 ounces and the 51 gauge Sample Film had a film weight/load of 9.0 ounces. The Paragon film had a containment force of 35 lbs compared with 55 lbs for the 63 gauge Sample Film and 50 lbs for the 51 gauge Sample Film.

#### **Comparative Example 27**

[0076] A Sample Film was prepared for comparison with a film manufactured by Pactiv under the tradename Max I. The Pactiv film was an 80 gauge film while the Sample Film was a 63 gauge film. It was determined that the Pactiv film had a film weight/load of 5.2 ounces and the Sample Film had a film weight/load of 4.2 ounces. The Pactiv film had a containment force of 22 lbs compared with 24 lbs for the Sample Film.

#### **Comparative Example 28**

[0077] A Sample Film was prepared for comparison with a film manufactured by AEP. The AEP film was an 80 gauge film while the Sample Film was a 51 gauge film. It was

determined that the AEP film had a film weight/load of 4.4 ounces and the Sample Film had a film weight/load of 3.0 ounces. The AEP film had a containment force of 12 lbs compared with 12 lbs for the Sample Film.

**[0078]** It will be obvious to those having skill in the art that many changes may be made to the details of the above-described embodiments of this invention without departing from the underlying principles thereof. The scope of the present invention should, therefore, be determined only by the following claims.